WEST Search History

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DATE: Saturday, June 26, 2004

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	DB=PGPB	B,USPT,USOC,EPAB,JPAB,DWPI; THES=ASSIGNEE; PLUR=YE	S; OP=ADJ
	L9	L8 and reduc\$3 near3 emission\$1	43
	L8	L7 and furnace	115
	L7	L6 and hydrogen near1 rich	224
	L6	naphtha near3 reform\$3 near4 hydrogen	533
	L5	L4 and reduc\$3 near4 carbon dioxide near3 emission\$1	3
Г	L4	L3 and naphtha with hydrogen	224
	L3	L2 and carbon dioxide	729
	L2	L1 and naphtha	994
	L1	(methane or natural gas) and partial oxidation	4591

END OF SEARCH HISTORY

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Search Results - Record(s) 1 through 5 of 5 returned.

☐ 1. Document ID: US 20030236312 A1

Using default format because multiple data bases are involved.

L15: Entry 1 of 5

File: PGPB

Dec 25, 2003

PGPUB-DOCUMENT-NUMBER: 20030236312

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030236312 A1

TITLE: Process for conversion of LPG and CH4 to syngas and higher valued products

PUBLICATION-DATE: December 25, 2003

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

O'Rear, Dennis J.

Petaluma

CA

US

US-CL-CURRENT: 518/728

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KWIC	Draw, De
		The special processor and the second		ANADORNI TODA PARAMENTO PRINCIPALIS AND A PARAMENTAL PROPERTY OF THE PARAMENT PROPERTY PROPERTY PROPERTY OF THE PA	og og til i i i araan i i i i i i i i ogg araab i i skrivis vangg	o#^^3***********************	001 x 80108013 x 301 \ 1.0 x 000 000 000 000 000 000 000 000 000		my diferentialism community and man		Programme Service Action	
	2. I	Documei	nt ID:	US 20	030191199	A1						
L15:	Entry	7 2 of	5				File:	PGPB		Oct	9,	2003

PGPUB-DOCUMENT-NUMBER: 20030191199

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030191199 A1

TITLE: REDUCTION OF CARBON DIOXIDE EMISSIONS FROM FISCHER-TROPSCH GTL FACILITY BY

AROMATICS PRODUCTION

PUBLICATION-DATE: October 9, 2003

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

O'Rear, Dennis J.

Petaluma

CA

US

APPL-NO: 10/ 118053 [PALM]
DATE FILED: April 9, 2002

INT-CL: [07] C07 C 27/06

US-CL-PUBLISHED: 518/705 US-CL-CURRENT: 518/705

REPRESENTATIVE-FIGURES: 1

ABSTRACT:

Provided is a process for reducing CO.sub.2 emissions generated by a Fischer-Tropsch GTL facility. The process includes introducing a synthesis gas into a Fischer-Tropsch reactor and performing a Fischer-Tropsch process to produce a Fischer-Tropsch product and CO.sub.2. At least a portion of the CO.sub.2 from the Fischer-Tropsch reactor is fed into at least one of a feed stream being fed to a synthesis gas formation reactor, producing the synthesis gas, or the synthesis gas being fed into the Fischer-Tropsch reactor. In addition, naphtha is obtained from the Fischer-Tropsch product and is fed into a naphtha reformer. Naphtha reforming is conducted, generating hydrogen by-product and C.sub.6-C.sub.10 product. At least a portion of the hydrogen by-product, generated during naphtha reforming, is fed into the feed stream, converting at least a portion of the CO.sub.2 in the feed stream into additional CO. Finally, the additional CO is converted into hydrocarbons in the Fischer-Tropsch reactor.

İ	Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KWIC	Draw, De
*******		01.00.1.00.0000000000000000000000000000				***************************************	hartuturusjät (1.65) armetuska	21/2011-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1		***************************************		romani i u u e ca ca u u u u u u u u u u u u u u u u u	97. TO 4700 77 X 27 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2

☐ 3. Document ID: US 20020120017 A1

L15: Entry 3 of 5

File: PGPB

Aug 29, 2002

PGPUB-DOCUMENT-NUMBER: 20020120017

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020120017 A1

TITLE: Processes for the production of hydrocarbons, power and carbon dioxide from carbon-containing materials

PUBLICATION-DATE: August 29, 2002

INVENTOR - INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Bohn, Mark S. Golden CO US

Benham, Charles B. Littleton CO US

APPL-NO: 09/ 963253 [PALM] DATE FILED: September 25, 2001

RELATED-US-APPL-DATA:

Application 09/963253 is a division-of US application 09/376709, filed August 17, 1999, US Patent No. 6306917

Application 09/376709 is a continuation-in-part-of US application 09/212374, filed December 16, 1998, ABANDONED

INT-CL: [07] C07 C 27/06

Record List Display Page 3 of 7

US-CL-PUBLISHED: 518/703 US-CL-CURRENT: 518/703

REPRESENTATIVE-FIGURES: 1

ABSTRACT:

Apparatus and processes for producing power, liquid hydrocarbons and carbon dioxide from heavy feedstocks, using a partial oxidation reactor to produce a synthesis gas, a Fischer-Tropsch reactor to convert the synthesis gas to hydrocarbon products and tail gases containing hydrogen and carbon dioxide, and a combined cycle plant to produce power from steam generated by recovering heat from the reactors and from combustible tail gases. By varying operating conditions and utilizing hydrogen for recycle to the Fischer-Tropsch reactor and/or hydrocracking wax products to produce lighter hydrocarbons, the process can selectively maximize the production of power, hydrocarbons or carbon dioxide. In preferred embodiments, the Fischer-Tropsch reactor can be a slurry reactor and can employ an iron-based catalyst.

CROSS REFERENCED PATENTS

[0001] This application is a divisional application of U.S. application Ser. No. 09/376,709 filed Aug. 17, 1999 and issued as U.S. Pat. No. ----- on ----- which is a continuation-in-part of U.S. application Ser. No. 09/212,374 filed Dec. 16, 1998.

Full	Title	e Citation Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KWIC	Draw. De
Military security		and in the control of	***************************************		or - 11 2 6 55 4000 - 3400 500 00 10 10 0		***************************************	······································		***************************************	
	4.	Document ID:	US 66	93138 B2							

File: USPT

Feb 17, 2004

US-PAT-NO: 6693138

L15: Entry 4 of 5

DOCUMENT-IDENTIFIER: US 6693138 B2

TITLE: Reduction of carbon dioxide emissions from Fischer-Tropsch GTL facility by aromatics production

DATE-ISSUED: February 17, 2004

INVENTOR - INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

O'Rear; Dennis J. Petaluma CA

ASSIGNEE-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY TYPE CODE

Chevron U.S.A. Inc. San Ramon CA 02

APPL-NO: 10/ 118053 [PALM]
DATE FILED: April 9, 2002

INT-CL: [07] C07 C 27/00, C10 G 35/00

US-CL-ISSUED: 518/700; 208/133, 208/141

Record List Display Page 4 of 7

US-CL-CURRENT: 518/700; 208/133, 208/141

FIELD-OF-SEARCH: 518/700, 208/133, 208/141

PRIOR-ART-DISCLOSED:

U.S. PATENT DOCUMENTS

PAT-NO	ISSUE-DATE	PATENTEE-NAME	US-CL
5620670	April 1997	Benham et al.	
6043288	March 2000	DeGeorge et al.	
6103773	August 2000	Wittenbrink et al.	

FOREIGN PATENT DOCUMENTS

FOREIGN-PAT-NO	PUBN-DATE	COUNTRY	US-CL
1006745	October 1965	GB	
01/72674	October 2001	WO	

OTHER PUBLICATIONS

U.S. Patent Application No. 10/118,029, Dennis J. O'Rear, Reducing CO2 levels in CO2-rich Natural Gases Converted into Liquid Fuels, Filed Apr. 9, 2002, attorney docket No. 005950-710.

United Kingdom Search Report dated Aug. 26, 2003.

ART-UNIT: 1621

PRIMARY-EXAMINER: Parsa; J.

ATTY-AGENT-FIRM: Burns, Doane, Swecker & Mathis, L.L.P.

ABSTRACT:

Provided is a process for reducing CO.sub.2 emissions generated by a Fischer-Tropsch GTL facility. The process includes introducing a synthesis gas into a Fischer-Tropsch reactor and performing a Fischer-Tropsch process to produce a Fischer-Tropsch product and CO.sub.2. At least a portion of the CO.sub.2 from the Fischer-Tropsch reactor is fed into at least one of a feed stream being fed to a synthesis gas formation reactor, producing the synthesis gas, or the synthesis gas being fed into the Fischer-Tropsch reactor. In addition, naphtha is obtained from the Fischer-Tropsch product and is fed into a naphtha reformer. Naphtha reforming is conducted, generating hydrogen by-product and C.sub.6 -C.sub.10 product. At least a portion of the hydrogen by-product, generated during naphtha reforming, is fed into the feed stream, converting at least a portion of the CO.sub.2 in the feed stream into additional CO. Finally, the additional CO is converted into hydrocarbons in the Fischer-Tropsch reactor.

17 Claims, 2 Drawing figures

Full Title	Citation	Front	Review	Classification	Date	Reference		Claims	KWIC	Drawt De

5. Document ID: US 6306917 B1

L15: Entry 5 of 5

File: USPT

Oct 23, 2001

US-PAT-NO: 6306917

DOCUMENT-IDENTIFIER: US 6306917 B1

TITLE: Processes for the production of hydrocarbons, power and carbon dioxide from carbon-containing materials

5

DATE-ISSUED: October 23, 2001

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Bohn; Mark S.

Golden CO

Benham; Charles S. Littleton CO

ASSIGNEE-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY TYPE CODE

Rentech, Inc. Denver CO 02

APPL-NO: 09/ 376709 [PALM] DATE FILED: August 17, 1999

PARENT-CASE:

CROSS REFERENCED PATENTS This application is a continuation-in-part of. U.S. application Ser. No. 09/212,374 filed Dec. 16, 1998 now abandoned.

INT-CL: [07] C07 C 27/00

US-CL-ISSUED: 518/700; 518/702, 518/703, 518/715, 518/721 US-CL-CURRENT: 518/700; 518/702, 518/703, 518/715, 518/721

FIELD-OF-SEARCH: 518/703, 518/702, 518/700, 518/721, 518/715

PRIOR-ART-DISCLOSED:

U.S. PATENT DOCUMENTS

PAT-NO	ISSUE-DATE	PATENTEE-NAME	US-CL
3617498	November 1971	Kittrell	
3972958	August 1976	Garwood et al.	
3986349	October 1976	Egan	
4059648	November 1977	Derr et al.	
4092825	June 1978	Egan	
4197184	April 1980	Munro et al.	
4256654	March 1981	Schlinger et al.	
4404088	September 1983	Bachtel et al.	
4433065	February 1984	Van Der Burgt et al.	518/703
4496371	January 1985	Urban et al.	

Record List Display Page 6 of 7

4501655	February 1985	Hilfman et al.	
4549396	October 1985	Garwood et al.	
4579985	April 1986	Minderhoud et al.	
4595702	June 1986	Chu et al.	
4605639	August 1986	Dyer et al.	
4617288	October 1986	Bell et al.	
4902303	February 1990	Den Bleyker	
4957715	September 1990	Grover et al.	
4992081	February 1991	Den Bleyker	
4994428	February 1991	Bell et al.	
5023276	June 1991	Yarrington et al.	
5026472	June 1991	Hoehn et al.	
5324335	June 1994	Benham et al.	
5324336	June 1994	Child	
5403568	April 1995	Stowe, Jr.	
5424051	June 1995	Nagji et al.	
5500449	March 1996	Benham et al.	
5504118	April 1996	Benham et al.	
5506272	April 1996	Benham et al.	
5543437	August 1996	Benham et al.	
5545238	August 1996	Brooker et al.	
5620670	April 1997	Benham et al.	
5621155	April 1997	Benham et al.	
5645613	July 1997	Benham et al.	
5666800	September 1997	Sorensen et al.	60/39.02
5763716	June 1998	Benham et al.	

OTHER PUBLICATIONS

Author: Marcel Dekker, Inc.; Title: The <u>Fischer-Tropsch</u> Synthesis in the Liquid Phase; Date: 1980; pp. 225-275.

Author: Chemical Engineering; Title: Technology to Cool Down Global Warming; Date: 1999; pp. 37,39,41.

Author: Alberto Rivalts; Title: Orimulsion. A new fuel for power generation and future feedstock use; Date: 1996; pp. 342-344.

Author: Charles B. Benham and Mark S. Bohn; Title: Maximization of Diesel Fuel Production from an Iron-Base Fischer-Tropsch Catalyst; Date: Dec. 1998; pp. 2-5. Author: A.R. Jones; Title: The commercial combustion of Orimulsion; Date: 1997; pp. 318-339.

Author: Mark S. Bohn and Charles B. Benham; Title: A Comparative Study of Alternate Flowsheets Using Orimusion as Feedstock; Date: Jan. 1999; pp. 1-10.

Author: David Gray and Glen Tomlinson; Title: A Novel Configuration for Coproducing Fischer-Tropsch Fuels and Electric Power from Coal and Natural Gas; Date: Sep. 1997; pp. 1-6.

Author: Dr. Benham and Dennis Yakobson; Title: Optimization of Conversion of Low Hydrogen Containing Feedstocks? Using Rentech's Gas-To-Liquids (GTL) Technology; Date: Jul. 1998; pp. 2-19.

Author: Schlesinge, Crowell, Leva, Storch; Title: Fischer-Tropsch Synthesis on Slurry Phase; Date: Jun. 1951; pp. 1474-1479.

Author: Arthur W. Tower III; Title: Gas-to-Liquids, Solids-yo-Liquids, Liquids-to-Liquids; Date: Dec. 18, 1998; Pages: whole book.

ART-UNIT: 161

PRIMARY-EXAMINER: Kumar; Shailendra

ASSISTANT-EXAMINER: Parsa; J.

ATTY-AGENT-FIRM: Martin; Rick Patent Law Offices of Rick Martin P.C.

ABSTRACT:

Apparatus and processes for producing power, liquid hydrocarbons and carbon dioxide from heavy feedstocks, using a partial oxidation reactor to produce a synthesis gas, a Fischer-Tropsch reactor to convert the synthesis gas to hydrocarbon products and tail gases containing hydrogen and carbon dioxide, and a combined cycle plant to produce power from steam generated by recovering heat from the reactors and from combustible tail gases. By varying operating conditions and utilizing hydrogen for recycle to the Fischer-Tropsch reactor and/or hydrocracking wax products to produce lighter hydrocarbons, the process can selectively maximize the production of power, hydrocarbons or carbon dioxide. In preferred embodiments, the Fischer-Tropsch reactor can be a slurry reactor and can employ an iron-based catalyst.

13 Claims, 5 Drawing figures

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Term	Documents
NAPHTHA	46317
NAPHTHAS	9911
HYDROGEN	980164
HYDROGENS	21638
FUEL	773287
FUELS	91272
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(L14 AND NAPHTHA WITH HYDROGEN SAME FUEL).PGPB,USPT,USOC,EPAB,JPAB,DWPI.	5

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FILE COVERS 1907 - 26 Jun 2004 VOL 141 ISS 1 FILE LAST UPDATED: 25 Jun 2004 (20040625/ED)

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=> methane and partial oxid?
METHANE IS NOT A RECOGNIZED COMMAND
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"HELP COMMANDS" at an arrow prompt (=>).

"HELP COMMANDS" at an arrow prompt (=>). => s (methane or natural gas or hydrocarbon) (1) partial oxid? 153496 METHANE 3204 METHANES 154844 METHANE (METHANE OR METHANES) 616643 NATURAL 31 NATURALS 616661 NATURAL (NATURAL OR NATURALS) 1364007 GAS 470209 GASES 1532769 GAS (GAS OR GASES) 64195 NATURAL GAS (NATURAL (W) GAS) 307933 HYDROCARBON 307105 HYDROCARBONS 473314 HYDROCARBON (HYDROCARBON OR HYDROCARBONS) 346013 PARTIAL 919 PARTIALS 346574 PARTIAL (PARTIAL OR PARTIALS) 2615075 OXID? 9536 PARTIAL OXID?

(PARTIAL(W)OXID?)

L1 3750 (METHANE OR NATURAL GAS OR HYDROCARBON) (L) PARTIAL OXID?

=> s l1 and synthesis gas 1126911 SYNTHESIS

3 SYNTHESISES

61904 SYNTHESES

1162021 SYNTHESIS

(SYNTHESIS OR SYNTHESISES OR SYNTHESES)

1364007 GAS 470209 GASES

1532769 GAS

1332,03 GAB

(GAS OR GASES) 14530 SYNTHESIS GAS

(SYNTHESIS (W) GAS)

L2 1434 L1 AND SYNTHESIS GAS

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=> s 12 and reform? (L) naphtha
        41708 REFORM?
        20398 NAPHTHA
         1652 NAPHTHAS
         20755 NAPHTHA
                 (NAPHTHA OR NAPHTHAS)
          3884 REFORM? (L) NAPHTHA
           13 L2 AND REFORM? (L) NAPHTHA
=> s 113 and carbon dioxide
L13 NOT FOUND
The L-number entered could not be found. To see the definition
of L-numbers, enter DISPLAY HISTORY at an arrow prompt (=>).
=> s 13 and carbon dioxide
      1061088 CARBON
        23608 CARBONS
       1069693 CARBON
                (CARBON OR CARBONS)
        412819 DIOXIDE
         6328 DIOXIDES
        414408 DIOXIDE
                (DIOXIDE OR DIOXIDES)
       192624 CARBON DIOXIDE
                (CARBON(W)DIOXIDE)
             3 L3 AND CARBON DIOXIDE
L4
=> d 14 ibib ab 1-3
    ANSWER 1 OF 3 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2004:203348 CAPLUS
                        140:220494
DOCUMENT NUMBER:
                        Dynamic sulfur tolerant process and system with inline
TITLE:
                        acid gas-selective removal for generating hydrogen for
                        fuel cells
                        Randhava, Surjit; Ho, W. S. Winston; Kao, Richard L.;
INVENTOR(S):
                        Camara, Elias H.
PATENT ASSIGNEE(S):
                        H2Fuel, LLC, USA; University of Kentucky Research
                        Foundation
SOURCE:
                        U.S. Pat. Appl. Publ., 19 pp.
                        CODEN: USXXCO
DOCUMENT TYPE:
                        Patent
                        English
LANGUAGE:
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:
     PATENT NO. KIND DATE APPLICATION NO. DATE
    US 2004047799 A1 20040311 US 2002-236324 US 2002-236324
                                       US 2002-236324 20020906
                                                          20020906
PRIORITY APPLN. INFO.:
    A dynamic sulfur tolerant process with inline acid gas-selective removal
     for converting sulfur-containing carbonaceous fuels into hydrogen or hydrogen
     rich gases includes the following steps: vaporizing liquid fuels and water
     in a vaporizer/preheater by transferring heat from reformate
    gas; feeding fuel, water and oxidant (air or oxygen) over a sulfur
     tolerant catalyst in an autothermal hydrodesulfurizing reformer
     (AHR) to yield a hydrogen rich reformate gas to combine the heat
    effects of the partial oxidn., steam reforming
    reactions, preheat and heat losses; converting CO and steam in the
    reformate gas to addnl. hydrogen and CO2 via a water gas shift
     (WGS) reaction in a WGS reactor; vaporizing and superheating water fed to
    a WGS boiler coil in a steam generator; removing selectively substantial
    amts. of acid gas (H2S and CO2) in WGS product gas in an acid-gas
```

selective membrane separator (ASMS); further reducing CO and CO2 concentration

levels in a methanator; and interchanging heat between hydrogen-rich gas and sweep gas in a heat interchanger. The ASMS product gas at 180-230° containing 0.5-2 mol% CO and <100 ppb H2S is an ideal feed for phosphoric acid fuel cells. The ASMS enables the methanation reaction equilibrium to decrease the CO and CO2 concns. in the hydrogen-rich gas to appropriate levels suitable for feeding PEM and alkaline fuel cells. The acid gas-selective membrane in the ASMS consists of polyamine and polyimide blends and copolymers which have a high acid gas/H2 selectivity and a good thermal stability of \leq 325°C. The membrane can consist of poly (2 or 4-vinylpyridine) and polyimide derived from polyamic acid which is synthesized from oxydianiline and oxydiphthalic anhydride. The fuel can be gasoline, diesel, naphtha, natural gas , liquefied petroleum gas (LPG), heating oil, kerosene, jet fuel, alkanes, alkenes, aroms., and hydrocarbon streams and alcs. selected from the group consisting of methanol, ethanol, butanol, pentanol, or alc.-containing steams. The AHR catalyst contains a Group IV rare earth metal sulfide, or a Group IV rare earth metal sulfate as hydrodesulfurization portion, a Group VIII transition metal and an alloy of a Group VIII transition metal as a dehydrogenation portion, and a ceramic oxide, a doped oxide, such as ZrO2, CeO2, Bi2O3, BiVO4, LaGdO3, Gd doped ceria, as an oxidation portion.

L4 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 2002:368370 CAPLUS

DOCUMENT NUMBER: 136:388285

TITLE: Method and catalyst for hydrocarbon reforming to

produce hydrogen and syngas

INVENTOR(S): Fukunaga, Tetsuya

PATENT ASSIGNEE(S): Idemitsu Kosan Co., Ltd., Japan

SOURCE: PCT Int. Appl., 27 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

```
APPLICATION NO. DATE
                  KIND DATE
     PATENT NO.
     WO 2002038268 A1 20020516 WO 2001-JP9660 20011105
         W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
             CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
             GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
             LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,
             PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA,
             UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
             DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,
             BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
     AU 2002011004 A5 20020521 AU 2002-11004 20011105
EP 1338335 A1 20030827 EP 2001-979007 20011105
         R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
     US 2004014600
                     A1 20040122
                                          US 2003-415558
                                                           20030501
PRIORITY APPLN. INFO.:
                                       JP 2000-339897 A 20001108
                                       WO 2001-JP9660 W 20011105
```

AB A catalyst suitable for use in various reforming reactions of hydrocarbons; and a process for producing H2 or synthesis gas which comprises using this catalyst to reform light hydrocarbons, are described. The catalyst comprises an alumina support containing Ce oxide and supported thereon (1) ≥1 Pt-group metals such as Ru, Pt, Rh, Pd, and Ir, (2) a Co and/or Ni ingredient, and (3) optionally an alkaline earth metal ingredient. The catalyst is useful for steam reforming, thermal self-reforming, partial oxidative reforming, or carbon dioxide

reforming of a hydrocarbon to produce H2 or synthesis

qas

REFERENCE COUNT:

THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

1993:606878 CAPLUS

DOCUMENT NUMBER:

119:206878

TITLE:

Process for the production of synthesis

gas

INVENTOR(S):

Hiles, Andrew George

PATENT ASSIGNEE(S):

Davy McKee (London) Ltd., UK

SOURCE:

PCT Int. Appl., 34 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE

WO 9315999 A1 19930819 WO 1993-GB329 19930217

W: AT, AU, BB, BG, BR, CA, CH, CZ, DE, DK, ES, FI, GB, HU, JP, KP, KR, LK, LU, MG, MN, MW, NL, NO, NZ, PL, PT, RO, RU, SD, SE, SK, UA, US

RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, SN, TD, TG

AU 9335083 A1 19930903 AU 1993-35083 19930217

PRIORITY APPLN. INFO:: GB 1992-3375 19920218

WO 1993-GB329 19930217

ΔR Synthesis gas is prepared from a hydrocarbon feedstock, e.g., natural gas, liquefied petroleum gas, naphtha, or a gas oil, by subjecting a part thereof to steam reforming and another part to partial oxidn. The hot gas stream from the steam reformer at 650-950° and 30-60 bar contains typically unconverted hydrocarbon and is mixed with the hot gas from the partial oxidn. unit which is at the same pressure as the hot gas from the steam reformer and at a temperature of 1200-1500°. The mixed stream at 1100-1200° can be passed through a secondary reforming zone to reduce CH4 content in the mixed stream to 2-5% or more on a dry gas basis. The exit gas from the secondary reformer can then be cooled and compressed for use, in MeOH synthesis. By adjusting the ratio of amts. of hydrocarbon feedstock fed to the steam reformer and to the partial oxidn. unit, the steam:C ratio in the steam reformer, and the O:C ratio for the partial oxidn. unit, the stoichiometry of the resulting synthesis gas can be adjusted to the desired value.

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=> s reduc? (2a) carbon dioxide emission?
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1836537 REDUC?

811690 REDN

43888 REDNS

839203 REDN

(REDN OR REDNS)

2299730 REDUC?

(REDUC? OR REDN)

1061088 CARBON

23608 CARBONS

1069693 CARBON

(CARBON OR CARBONS)

412819 DIOXIDE

6328 DIOXIDES

```
414408 DIOXIDE
                (DIOXIDE OR DIOXIDES)
        470318 EMISSION?
          1488 CARBON DIOXIDE EMISSION?
                 (CARBON (W) DIOXIDE (W) EMISSION?)
           316 REDUC? (2A) CARBON DIOXIDE EMISSION?
L5
=> s 15 and gas (1a) liquid
       1364007 GAS
        470209 GASES
       1532769 GAS
                (GAS OR GASES)
        654512 LIQUID
        119450 LIQUIDS
        744941 LIQUID
                (LIQUID OR LIQUIDS)
        900719 LIQ
         85888 LIQS
        934006 LIQ
                 (LIQ OR LIQS)
       1303653 LIQUID
                 (LIQUID OR LIQ)
         81873 GAS (1A) LIQUID
            3 L5 AND GAS (1A) LIQUID
1.6
=> s 15 and naphtha (2a) reform?
         20398 NAPHTHA
          1652 NAPHTHAS
         20755 NAPHTHA
                 (NAPHTHA OR NAPHTHAS)
         41708 REFORM?
          3143 NAPHTHA (2A) REFORM?
L7
             1 L5 AND NAPHTHA (2A) REFORM?
=> s 16 or 17
            3 L6 OR L7
=> d 18 ibib ab 1-3
    ANSWER 1 OF 3 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2003:796335 CAPLUS
                         139:293676
DOCUMENT NUMBER:
                         Reduction of carbon
TITLE:
                         dioxide emissions from a
                         Fischer-Tropsch gas-to-liquid
                         facility by aromatics production
INVENTOR(S):
                         O'Rear, Dennis J.
                         Chevron U.S.A., Inc., USA U.S. Pat. Appl. Publ., 9 pp.
PATENT ASSIGNEE(S):
SOURCE:
                         CODEN: USXXCO
DOCUMENT TYPE:
                         Patent
LANGUAGE:
                         English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:
                                           APPLICATION NO. DATE
     PATENT NO.
                KIND DATE
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|----|------|-------|-----|-----|-----|------|------|-----|-----|------|-------|-------|-----|------|------|-----|-----|
| US | 2003 | 1911 | 99 | A | 1 | 2003 | 1009 | | U | 3 20 | 02-1 | 18053 | 3 | 2002 | 0409 | | |
| US | 6693 | 138 | | B2 | 2 | 2004 | 0217 | | | | | | | | | | |
| WO | 2003 | 08726 | 54 | A2 | 2 | 2003 | 1023 | | W | 200 | 3 - U | S875 | 9 | 2003 | 0320 | | |
| WO | 2003 | 08726 | 54 | A3 | 3 . | 2003 | 1224 | | | | | | | | | | |
| | W: | ΑE, | AG, | AL, | AM, | ΑT, | AU, | ΑZ, | BA, | BB, | BG, | BR, | BY, | ΒZ, | CA, | CH, | CN, |
| | | CO, | CR, | CU, | CZ, | DE, | DK, | DM, | DZ, | EC, | EE, | ES, | FΙ, | GB, | GD, | GE, | GH, |
| | | GM, | HR, | HU, | ID, | IL, | IN, | IS, | JP, | KE, | KG, | KP, | KR, | KΖ, | LC, | LK, | LR, |

LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG GB 2003-7419 GB 2388117 A1 20031105 20030331 20031013 NL 2003-1023136 20030409 NL 1023136 Α1 US 2002-118053 A 20020409 PRIORITY APPLN. INFO.: The process includes introducing a synthesis gas into a Fischer-Tropsch reactor and performing a Fischer-Tropsch process to produce a Fischer-Tropsch product and CO2 . A portion of the CO2 from the Fischer-Tropsch reactor is fed into at least one of a feed stream being fed to a synthesis gas formation reactor, producing the synthesis gas, or the synthesis gas being fed into the Fischer-Tropsch reactor. Naphtha is obtained from the Fischer-Tropsch product and is fed into a naphtha reformer generating a hydrogen byproduct and a C6-10 product. A portion of the hydrogen byproduct, generated during naphtha reforming, is fed into the feed stream, converting a portion of the CO 2 in the feed stream into addnl. CO and the addnl. CO is converted into hydrocarbons in the Fischer-Tropsch reactor.

L8 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2002:385538 CAPLUS

DOCUMENT NUMBER:

137:65455

TITLE:

CO2 mitigation and fuel production

AUTHOR(S):

Steinberg, M.

CORPORATE SOURCE:

Department of Applied Science, Brookhaven National

Laboratory, Upton, NY, 11973, USA

SOURCE:

ACS Symposium Series (2002), 809(CO2 Conversion and

Utilization), 31-38

CODEN: ACSMC8; ISSN: 0097-6156 American Chemical Society

PUBLISHER:

American Chemica Journal

DOCUMENT TYPE: LANGUAGE:

English

CO2 mitigation technologies deals with how to utilize fossil fuels, coal, oil and gas with reduced CO2 emissions. Most development work to date has emphasized improving efficiency in generation and utilization of energy and in removal and recovery of CO2 from central power stations followed by disposal in underground wells or in the ocean. The latter suffers from economic penalties and potential adverse environmental effects. CO2 utilization for the chems. industry is problematic because of the capacity mismatch between the gross CO2 emission and the relatively smaller chemical products market. However, utilization of CO2 for conversion to alternative fuels for stationary and automotive power has potential of matching capacity between emissions and utilization. Although hydrogen-rich gaseous fuels such as methane and hydrogen can be used as alternative automotive liquid fuels which includes methanol and other higher oxygenates appear safer and fit in with the current liquid fuel infrastructure. The key for gas to liq. conversion utilizing CO2 is the production of hydrogen for conversion with CO2. Carnol process, catalytically reacts CO2 from coal fired plants with hydrogen from the thermal decomposition of methane while sequestering the elemental carbon, to produce methanol and higher oxygenated fuels. CO2 emission redns. approaching 80% using fuel cell engines can be achieved compared to the conventional system of coal-fired power generating plants and gasoline-driven IC automotive engines.

REFERENCE COUNT:

THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2004 ACS on STN ACCESSION NUMBER: 1993:606860 CAPLUS

DOCUMENT NUMBER:

119:206860

TITLE:

Evaluation of liquid fuel production technology from

natural gas and coal from the viewpoint of the

reduction of carbon dioxide

emission

AUTHOR(S):

Inaba, Atsushi; Saito, Ikuo

CORPORATE SOURCE: SOURCE:

Natl. Inst. Resour. Environ., Tsukuba, Japan Nippon Eneruqi Gakkaishi (1993), 72(6), 457-66

CODEN: NENGEM; ISSN: 0916-8753

DOCUMENT TYPE:

Journal Japanese

LANGUAGE:

Technologies for production of electricity and liquid fuel from natural gas and/or coal were evaluated from the viewpoint of reduction of CO2 emissions in the long run, when petroleum would not be produced. If natural gas supply are not sufficient to satisfy demands of both electricity and liquid fuel production and if those technologies are not improved in their thermal efficiency from the present practical levels, natural gas used for the power generation should be replaced with coal because CO2 emissions from such systems would be smaller than by systems in which coal is used for liquid fuel production However, if direct coal liquefaction technol. is improved near the level of theor. reaction, the utilization of coal for

In

this case, if hydrogen produced from natural gas is used for direct coal liquefaction, and CO2 emissions can be reduced much more. In integrated energy systems, in which excess hydrogen from liquid fuel production from natural gas is used for the indirect liquefaction of coal, is useful for the reduction of CO2 emissions when the reaction heat of steam reforming of natural gas is supplied from the outside such as high-temperature gas-cooled nuclear reactor.

liquid fuel production instead of power generation can reduce CO2 emissions.